

Europäisches Patentamt  
European Patent Office  
Office européen des brevets



(11) **EP 1 329 230 A1**

(12) **EUROPEAN PATENT APPLICATION**

(43) Date of publication:  
23.07.2003 Bulletin 2003/30

(51) Int Cl.7: **A61L 31/10**, B05D 1/00,  
G03F 7/16

(21) Application number: **03000600.1**

(22) Date of filing: **14.01.2003**

(84) Designated Contracting States:  
**AT BE BG CH CY CZ DE DK EE ES FI FR GB GR**  
**HU IE IT LI LU MC NL PT SE SI SK TR**  
Designated Extension States:  
**AL LT LV MK RO**

(72) Inventor: **Sundar, Rangarajan**  
**Santa Rosa, CA 95404 (US)**

(74) Representative: **Hughes, Andrea Michelle**  
**Frank B. Dehn & Co.,**  
**European Patent Attorneys,**  
**179 Queen Victoria Street**  
**London EC4V 4EL (GB)**

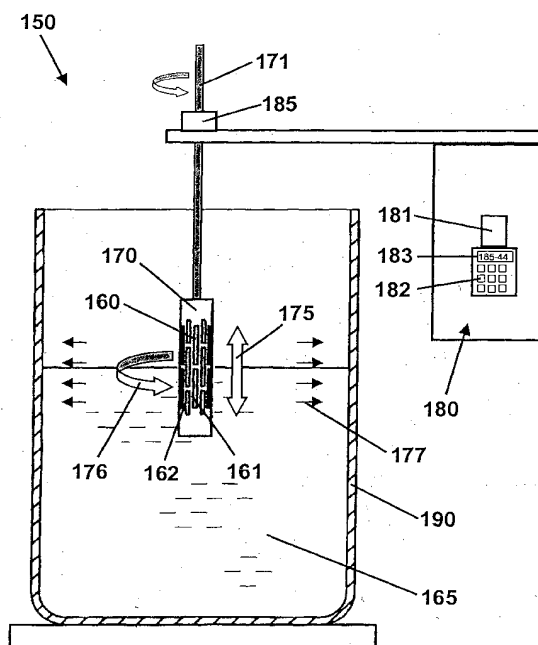
(30) Priority: **14.01.2002 US 50219**

(71) Applicant: **Medtronic AVE, Inc.**  
**Santa Rosa, California 95403 (US)**

(54) **Stent delivery system, device, and method for coating**

(57) The invention provides a stent delivery system, a stent device, and a method for coating a stent. The stent delivery system includes a catheter, a balloon operably attached to the catheter, and a stent disposed on the balloon. The stent includes at least one coating applied by dipping a portion of the stent into a coating liquid while simultaneously rotating the stent. The stent device

includes a body and at least one coating rotationally applied to a portion of the body, while the body is at least partially immersed in a coating liquid. The stent coating method includes immersing a portion of the stent into a coating liquid, and withdrawing the immersed portion of the stent from the coating liquid. The stent is simultaneously rotated with respect to the coating liquid while the stent is being immersed and withdrawn.



**FIG. 4**

## Description

**[0001]** The present invention relates generally to the field of implantable medical devices. More particularly, the invention relates to a stent delivery system and device having a coating, and a method for coating the same.

**[0002]** Balloon angioplasty has been used for the treatment of narrowed and occluded blood vessels. A frequent complication associated with the procedure is restenosis, or vessel re-narrowing. Within 3-6 months of angioplasty, restenosis occurs in almost 50 percent of patients. In order to reduce the incidence of re-narrowing, several strategies have been developed. Implantable devices, such as stents, have been used to reduce the rate of angioplasty related restenosis by about half. The use of such devices has greatly improved the prognosis of these patients. Nevertheless, restenosis remains a formidable problem associated with the treatment of narrowed blood vessels.

**[0003]** Restenosis associated with interventional procedures such as balloon angioplasty may occur by two mechanisms: thrombosis and intimal hyperplasia. During angioplasty, a balloon is inflated within an affected vessel thereby compressing the blockage and imparting a significant force, and subsequent trauma, upon the vessel wall. The natural antithrombogenic lining of the vessel lumen may become damaged thereby exposing thrombogenic cellular components, such as matrix proteins. The cellular components, along with the generally antithrombogenic nature of any implanted materials (e. g., a stent), may lead to the formation of a thrombus, or blood clot. The risk of thrombosis is generally greatest immediately after the angioplasty.

**[0004]** The second mechanism of restenosis is intimal hyperplasia, or excessive tissue re-growth. The trauma imparted upon the vessel wall from the angioplasty is generally believed to be an important factor contributing to hyperplasia. This exuberant cellular growth may lead to vessel "scarring" and significant restenosis. The risk of hyperplasia associated restenosis is usually greatest 3 to 6 months after the procedure.

**[0005]** Prosthetic devices, such as stents or grafts, may be implanted during interventional procedures such as balloon angioplasty to reduce the incidence of vessel restenosis. To improve device effectiveness, stents may be coated with one or more therapeutic agents providing a mode of localized drug delivery. The therapeutic agents are typically intended to limit or prevent the aforementioned mechanisms of restenosis. For example, antithrombogenic agents such as heparin or clotting cascade IIb/IIIa inhibitors (e.g., abciximab and eptifibatide) may be coated on the stent thereby diminishing thrombus formation. Such agents may effectively limit clot formation at or near the implanted device. Some antithrombogenic agents, however, may not be effective against intimal hyperplasia. Therefore, the stent may also be coated with antiproliferative agents or

other compounds to reduce excessive endothelial re-growth. Therapeutic agents provided as coatings on implantable medical devices may effectively limit restenosis and reduce the need for repeated treatments.

**[0006]** Several considerations should be made when devising a strategy for coating implantable prosthetic devices, such as stents or grafts. One consideration in coating strategy relates to surface uniformity. Ideally, coatings should be evenly applied with limited surface imperfections. Some coating strategies, however, may produce pooling of the coating material and/or dry spots. Failure to control surface uniformity may lead to inaccurate, non-uniform drug dose delivery and therapeutic variability from device to device. Therefore, it would be desirable to provide a uniform stent coating.

**[0007]** Another consideration in coating strategy relates to topography. It may be desirable for the coating to be disposed on certain areas of the stent. For example, stents typically have a wire mesh with open spaces formed between. Some coating strategies may leave 'bridged' material within the open spaces. Such coating bridges may break off causing complications or may prevent the device from expanding or functioning properly. As another example, only certain portions of the stent may require coating, or several coating layers may be required. As such, it would be desirable to control the stent coating topography.

**[0008]** Another consideration in coating strategy relates to efficiency. It may be desirable to effectively coat the implantable device in relatively short time, with a minimal amount of coating material. Some coating strategies require lengthy steps, thereby reducing the amount of devices that can be coated within a certain period. In addition, some strategies do not utilize coating material in a complete manner thereby increasing cost. For example, coating material that is vaporized may get dispersed on areas other than the stent surface. Therefore, it would be desirable to efficiently coat the stent.

**[0009]** Accordingly, it would be desirable to provide a strategy for coating a stent that would overcome the aforementioned and other disadvantages.

## SUMMARY OF THE INVENTION

**[0010]** According to one aspect, the invention provides a method for coating a stent comprising:

immersing a portion of the stent into a coating liquid; and  
withdrawing the immersed portion of the stent from the coating liquid; characterised by  
simultaneously rotating the stent with respect to the coating liquid while the stent is being immersed and withdrawn.

**[0011]** According to another aspect, there is provided a method for coating a stent comprising:

immersing a portion of the stent into a coating liquid; and characterised by simultaneously rotating the stent with respect to the coating liquid while the stent is at least partially immersed in the coating liquid.

**[0012]** The coating may include a therapeutic agent and may be substantially on an outer surface of the stent. The coating may be about 1 to 150 microns thick. The stent may be dipped at a rate of about 0.1 to 25.0 millimeters per second, and for a time period of about 5 seconds to 10 minutes. The stent may be rotated at a rate of about 100 to 25,000 rotations per minute. The system may further include a control sequence, and a programmable logic chip. The logic chip may control at least one of the dipping and rotation of the stent based on the control sequence.

**[0013]** Another aspect of the invention provides a stent coating device comprising:

means for immersing a portion of the stent into a coating liquid; and  
means for withdrawing the immersed portion of the stent from the coating liquid; and characterised by means for simultaneously rotating the stent with respect to the coating liquid while the stent is being immersed and withdrawn.

**[0014]** The coating may include a therapeutic agent and may be substantially on an outer surface of the body. The coating may be about 1 to 150 microns thick. The body may be dipped at a rate of about 0.1 to 25.0 millimeters per second, and for a time period of about 5 seconds to 10 minutes. The body may be rotated at a rate of about 100 to 25,000 rotations per minute. The device may further include a control sequence, and a programmable logic chip. The logic chip may control the rotational application based on the control sequence.

**[0015]** Another aspect of the invention provides a stent coating device comprising:

means for immersing a portion of the stent into a coating liquid; and characterised by  
means for simultaneously rotating the stent with respect to the coating liquid while the stent is at least partially immersed in said coating liquid.

**[0016]** The foregoing and other features and advantages of the invention will become further apparent from the following detailed description of the presently preferred embodiments, given by way of example only, read in conjunction with the accompanying drawings. The detailed description and drawings are merely illustrative of the invention, rather than limiting the scope of the invention being defined by the appended claims.

## BRIEF DESCRIPTION OF THE DRAWINGS

### [0017]

**FIG. 1** is a perspective view of a stent delivery system made in accordance with the present invention; **FIG. 2** is a perspective view of a prior art stent compatible with the coating process according to the present invention;

**FIG. 3** is a magnified view of two W-shaped elements of the stent in **FIG. 2**; and

**FIG. 4** is diagram of a stent coating process made in accordance with the present invention.

**[0018]** Referring to the drawings, wherein like reference numerals refer to like elements, **FIG. 1** is a perspective view of a stent delivery system made in accordance with the present invention and shown generally by numeral **100**. The stent delivery system **100** includes a catheter **105**, a balloon **110** operably attached to the catheter **105**, and a stent **120** disposed on the balloon **110**. The balloon **110**, shown in a collapsed state, may be any variety of balloons capable of expanding the stent **120**. The balloon **110** may be manufactured from any sufficiently elastic material such as polyethylene, polyethylene terephthalate (PET), nylon, or the like. In one embodiment, the balloon **110** may include retention means **111**, such as mechanical or adhesive structures, for retaining the stent **120** until it is deployed. The catheter **105** may be any variety of balloon catheters, such as a PTCA balloon catheter, capable of supporting a balloon during angioplasty.

**[0019]** The stent **120** may be any variety of implantable prosthetic devices capable of carrying a coating known in the art. In one embodiment, the stent **120** may have a plurality of identical cylindrical stent segments placed end to end. Four stent segments **121**, **122**, **123**, and **124** are shown, and it will be recognized by those skilled in the art that an alternative number of stent segments may be used. The stent **120** includes at least one coating applied by dipping a portion of the stent **120** into a coating liquid while simultaneously rotating the stent **120**. In one embodiment, three coating layers **125**, **126**, and **127** may be applied by a dip-spin coating process on various segments **121**, **122**, and **124**. Segment **121** is shown having two coating layers **125**, and **126**. Segment **122** and segment **124** are shown each having one coating layer **125**, and **127**, respectively. Segment **123** is shown having no coating. The coating layers **125**, **126**, and **127** are merely exemplary, and it should be recognized that other coating configurations are possible.

**[0020]** To describe the dip-spin coating process of the present invention, the following figures and description are provided. **FIG. 2** is a perspective view of a prior art stent **130** compatible with the coating process of the present invention. Those skilled in the art will recognize that numerous stents, grafts, and implantable prosthetic



devices are compatible with the disclosed coating method and that the described stent **130** example is merely one illustration of the process. The stent **130** is an example of a wire-tubular hybrid stent disclosed by U.S. Patent No. 5,935,162 issued to Dang.

[0021] The stent **130** includes a generally tubular body defining a passageway extending along a longitudinal axis **131**. The stent **130** is formed from a plurality of cylindrical segments **132** arranged successively along the longitudinal axis **131**. Each of cylindrical segments **132** has a length along the longitudinal axis **131** and includes a plurality of W-shaped elements **135**. The W-shaped elements **135** open in alternating directions along the longitudinal axis **131** about the perimeter or circumference of the cylindrical segments **132**. The W-shaped elements **135** are connected to each other by a tie member **136** that is attached to center sections of each of the W-shaped elements **135**.

[0022] The stent **130** is shown in an expanded state in which the cylindrical segments **132** have been expanded radially outward from the longitudinal axis **131**. The stent **130** may be compressed into a smaller diameter for delivery within a vessel lumen at which point the stent **130** may be expanded to provide support to the vessel. The stent **130** may be of the self-expanding variety and manufactured from nickel titanium alloys and other alloys that exhibit superelastic behavior (i.e., capable of significant distortion without plastic deformation). Alternatively, the stent **130** may be designed to be expanded by a balloon or some other device, and may be manufactured from an inert, biocompatible material with high corrosion resistance. The biocompatible material should ideally be plastically deformed at low-moderate stress levels. Suitable materials include, but are not limited to, tantalum, MP35N, stainless steel, titanium ASTM F63-83 Grade 1, niobium or high carat gold K 19-22.

[0023] FIG. 3 is a magnified view of two W-shaped elements **135** of the stent **130** in FIG. 1. The surface of the W-shaped elements is shown with a polymeric coating **140** having coating anomalies **141**. The anomalies **141** may take on numerous configurations, but common examples include pooling **142**, dry spots **143**, and bridging **144**. The anomalies **141** are typically produced during an imperfect coating process. Ideally, a coating process should produce an even coating **140** free of pooling **142**, dry spots **143**, and bridging **144**. Such coating anomalies **141** may lead to inaccurate, non-uniform drug dose delivery and therapeutic variability. In addition, anomalies such as bridging **144** may break off causing complications or may prevent the stent **130** from expanding or functioning properly. For example, bridging **144** materials may enter the bloodstream thereby releasing therapeutic agents in undesired locations. Accordingly, it is desirable to minimize coating anomalies **141**.

[0024] Turning now to FIG. 4, a stent coating process made in accordance with the present invention is shown

generally as numeral **150**. The coating process **150** provides one or more coatings on a portion of a stent device **160**. A stent device **160**, referred herein as  stent , may be any number of implantable prosthetic devices capable of carrying a coating. The stent **160** may include a body **161** and at least one coating **162** rotationally applied to a portion of the body **161**, while the body **161** is at least partially immersed in a coating liquid **165**. The stent **160** may be manufactured from a skeletal framework or mesh of material forming a tubelike structure and may be capable of self-expanding or being expanded by another device such as a balloon. The stent **160** material may include any number of metallic and polymeric biocompatible materials recognized in the art for such devices.

[0025] To apply the coating **162**, the stent **160** may be slidably mounted on a cylindrically shaped mandrel **170** that is fixably attached to a rod **171**. The stent **160**, mandrel **170**, and rod **171**, may be moved during the coating process **150** by an actuator **180**. The actuator **180** may provide vertical **175** (e.g., immersion and withdrawal) and rotational **176** movements. The actuator **180** may include a logic chip **181** programmed with a control sequence to control timing, speed, directionality, and variations of the vertical **175** and rotational **176** movement characteristics. The actuator **180** may further include a keypad **182** and display **183** for modifying and viewing portions of the control sequence, respectively. The control sequence and corresponding movement characteristics may be actuated by one or more motors **185** controlled by the logic chip **181**. The motors **185** may be of any variety of electric motors, hydraulic units, and the like, recognized in the art for providing rotational and linear movements.

[0026] The stent **160** may be dipped in the coating liquid **165** carried within a compatible container **190** (i.e., inert with respect to the coating liquid **165**). In one embodiment, the coating liquid **165** may be a polymeric solution. The polymeric solution may include one or more polymers, solvents, and therapeutic agents. The polymer may be dissolved in an appropriate solvent to provide a matrix for incorporating the therapeutic agent within the coating. Suitable polymers include, but are not limited to, urethane, polycaprolactone (PCL), polymethylmethacrylate (PMMA), and the like. Suitable solvents include, but are not limited to, acetone, ethyl acetate, tetrahydrofuran (THF), chloroform, N-methylpyrrolidone (NMP), and the like. Suitable therapeutic agents include, but are not limited to, antiangiogenesis agents, antiendothelin agents, antimetogenic factors, antioxidants, antiplatelet agents, antiproliferative agents, antisense oligonucleotides, antithrombogenic agents, calcium channel blockers, clot dissolving enzymes, growth factors, growth factor inhibitors, nitrates, nitric oxide releasing agents, vasodilators, virus-mediated gene transfer agents, agents having a desirable therapeutic application, and the like. Specific example of therapeutic agents include abciximab, angiopeptin, colchicine, ep-

tifibatide, heparin, hirudin, lovastatin, methotrexate, streptokinase, taxol, ticlopidine, tissue plasminogen activator, trapidil, urokinase, and growth factors VEGF, TGF-beta, IGF, PDGF, and FGF.

**[0027]** The stent coating process **150** may begin by preparing the coating liquid **165**, such as a polymeric solution. In one embodiment, preparation of the polymeric solution may include dissolving polymer in solvent. The choice of solvent controls the ability to place the polymer into solution and, thus, should be considered. The use of solvent blends may also be considered. A blend of solvents may be preferred to ensure uniform coating **162** with a smooth surface. Solvent volatility may also be considered. Too rapid solvent evaporation may result in surface imperfections; too slow solvent evaporation may result in handling defects as well as high solvent retention. As such, the polymer-solvent solution should ideally provide a uniform coating **162**, smooth surface without major imperfections, and little to no solvent retention. In one embodiment, the polymer-solvent solution may have a solids range from about 1 to 10 percent and a viscosity of about 5 to 30 centipoises.

**[0028]** One or more therapeutic agents may be added to the polymeric solution. Ideally, the polymer-solvent solution should be capable of dispersing the agent evenly throughout the solution. Furthermore, the polymer-solvent solution should not chemically alter the therapeutic character of the agent. Examples of suitable polymer/solvent/therapeutic agent combinations include: polylactic acid/trichloroethane/colchicines; polyurethane/THF/taxol; (poly(D,L-lactide)/PCL)/dimethylformamide/hirudin; (poly(D,L-lactide)/polyglycolide)/ethylacetate/ticlopidine; and polyethylene oxide/ethanol/heparin. These combinations are merely exemplary, and it should be recognized that other combinations are possible.

**[0029]** After the polymer and therapeutic agent have been dissolved in the solvent, the polymeric solution may be aged and cleared. In one embodiment, the solution may be mixed for 12 or more hours while being stirred to ensure complete dissolution and homogeneity. The solution may also be filtered with an appropriate filter to remove any impurities and haziness. Those skilled in the art will recognize that the preparation of the coating liquid **165** may vary greatly and may be contingent upon the materials used. After the coating liquid **165** has been prepared, it may be transferred to the container **190**.

**[0030]** The stent **160** mounted on the mandrel **170** may begin rotation as effected by the actuator motor **185**. The stent **160** may be rotated initially at a rate of about 100 to 3,500 rotations per minute (rpm). More preferably, the stent **160** may be rotated at about 500 to 700 rpm. The stent **160** may then be immersed in the coating liquid **165** at a rate of about 0.1 to 25.0 millimeters per second (mm/sec). More preferably, the stent **160** may be immersed at a rate of about 0.5 to 5.0 mil-

limeters per second. The immersion may continue until the stent body **161** portion requiring coating has been immersed. The stent **160** may be immersed for a total time of about 5 seconds to 10 minutes. More preferably, the stent **160** may be immersed for about 5 to 20 seconds. The immersion time may control a stent wetting characteristic; the characteristic relating to how completely the stent is covered with the coating liquid **165**. Ideally, the stent **160** should be evenly wetted free from any dry spots. A lower viscosity coating liquid **165** generally require shorter immersion time. The temperature of the coating liquid **165** affects its viscosity and, therefore, may also be a factor in determining immersion time.

**[0031]** Once the stent **160** has been immersed, the stent **160** may begin withdrawal wherein it is rotated at about 600 to 25,000 rpm. More preferably, the stent **160** may be rotated during withdrawal at a rate of about 3,000 to 10,000 rpm. The rate of rotation during withdrawal may influence the uniformity of the coating **162**. For example, too slow rotation during withdrawal may lead to coating pooling; too fast rotation may excessively remove coating material. The stent **160** may be withdrawn at a rate of about 0.1 to 25.0 mm/sec. More preferably, the stent **160** may be withdrawn initially at 0.1 mm/sec and then at 0.7mm/sec providing a uniform coating **162**. The withdrawal rate may control a coating thickness. Generally, faster withdrawal rates produce greater coating thickness. The coating **162** may be in the range of about 1 to 150 microns thick. More preferably, the coating **162** may be about 5 to 30 microns thick. The rotating stent **160** may experience a significant centrifugal force **177**. The coating liquid **165** may be forced substantially from an inner to an outer surface portion of the stent **160**. As a result, the bulk of the coating **162** and integral therapeutic agent may be positioned proximate to where the stent **160** contacts the vessel wall. This topography may provide efficient coating liquid **165** utilization and an effective drug delivery method.

**[0032]** Another factor affecting the coating process **150** pertains to the manner in which the coated stent **160** is dried. Ideally, the coated stent **160** should be dried to allow for a desirable rate of solvent evaporation. For example, too rapid drying causes moisture to be entrained in the coating **162** causing surface blemishes; too slow drying increases solvent retention. The drying conditions may vary greatly and are generally dependant upon the nature of the coating liquid **165** and, to a great extent, the solvent used. After the coating **162** has dried, the stent coating process **150** may be repeated to provide additional coating(s) positioned on top of, or on other locations of the stent **160**.

**[0033]** The described stent coating process **150** may provide a coating **162** that is controllable, uniform, efficient, and free from bridging and pooling anomalies. The process **150** may be optimized by customizing the timing, speed, directionality, and variations of the vertical **61** and rotational **62** movement characteristics. These

characteristics may be adjusted based on the nature of the stent **160** and coating liquid **165** used. The concurrent spinning and dipping of the stent **160** may speed the rate at which the coating **162** is applied. In addition, the process **150** may provide efficient utilization of coating liquid **165**. Accordingly, the process **150** may reduce the time and cost of conventional coating strategies.

**[0034]** While the embodiments of the invention disclosed herein are presently considered to be preferred, various changes and modifications can be made without departing from the spirit and scope of the invention. For example, the stent configuration is not limited to any particular stent design. In addition, the coating liquid composition and coating process movement characteristics may be varied considerably while providing a desirable coating. Upon reading the specification and reviewing the drawings hereof, it will become immediately obvious to those skilled in the art that myriad other embodiments of the present invention are possible, and that such embodiments are contemplated and fall within the scope of the presently claimed invention to the extent that this is defined in the appended claims.

## Claims

### 1. A method for coating a stent (120) comprising:

immersing a portion of the stent (120) into a coating liquid; and  
withdrawing the immersed portion of the stent from the coating liquid; **characterised by**  
simultaneously rotating the stent (120) with respect to the coating liquid while the stent is being immersed and withdrawn.

### 2. A method for coating a stent (120) comprising:

immersing a portion of the stent (120) into a coating liquid; and **characterised by** simultaneously rotating the stent with respect to the coating liquid while the stent is at least partially immersed in the coating liquid.

### 3. The method of claim 1 or 2 wherein the rotation forces the coating liquid to an outer portion of the stent.

### 4. The method of claim 1, 2 or 3 further comprising applying multiple layered coatings (125, 126, 127).

### 5. The method of any preceding claim 17 wherein immersing the stent (120) comprises controlling a stent wetting characteristic.

### 6. The method of any preceding claim wherein the stent (120) is immersed at a rate of 0.1 to 25.0 millimeters per second.

### 7. The method of any preceding claim wherein the stent (120) is immersed for a time period of about 5 seconds to 10 minutes.

### 8. The method of any preceding claim wherein the stent (120) is rotated during immersion at a rate of 100 to 3,500 rotations per minute.

### 9. The method of claim 1 or any claim dependent thereon wherein withdrawing the stent (120) comprises controlling a stent coating thickness.

### 10. The method of claim 9 wherein the stent (120) coating thickness comprises a thickness of 1 to 150 microns.

### 11. The method of claim 1 or any claim dependent thereon wherein the stent is withdrawn at a rate of 0.1 to 25.0 millimeters per second.

### 12. The method of claim 1 or any claim dependent thereon wherein the stent (120) is rotated during withdrawal at a rate of 600 to 25,000 rotations per minute.

### 13. The method any preceding claim further comprising:

programming a control sequence; and  
controlling at least one of the immersion, withdrawal, and rotation based on the control sequence.

### 14. The method of any preceding claim wherein the coating includes a therapeutic agent.

### 15. The method of any preceding claim wherein the coating is substantially on an outer surface of the stent.

### 16. A stent coating device comprising:

means (180) for immersing a portion of the stent (120, 160) into a coating liquid; and  
means (180) for withdrawing the immersed portion of the stent from the coating liquid; and

#### **characterised by**

means (180) for simultaneously rotating the stent (160) with respect to the coating liquid (165) while the stent is being immersed and withdrawn.

### 17. The device of claim 14 further comprising:

means (180) programmed with a control sequence; and  
means (180) for controlling at least one of the immersion, withdrawal, and rotation based on

the control sequence.

**18. A stent coating device comprising:**

means (180) for immersing a portion of the  
stent (120, 160) into a coating liquid; and

**characterised by**

means (180) for simultaneously rotating the  
stent with respect to the coating liquid (165) while  
the stent is at least partially immersed in said coat-  
ing liquid.

15

20

25

30

35

40

45

50

55

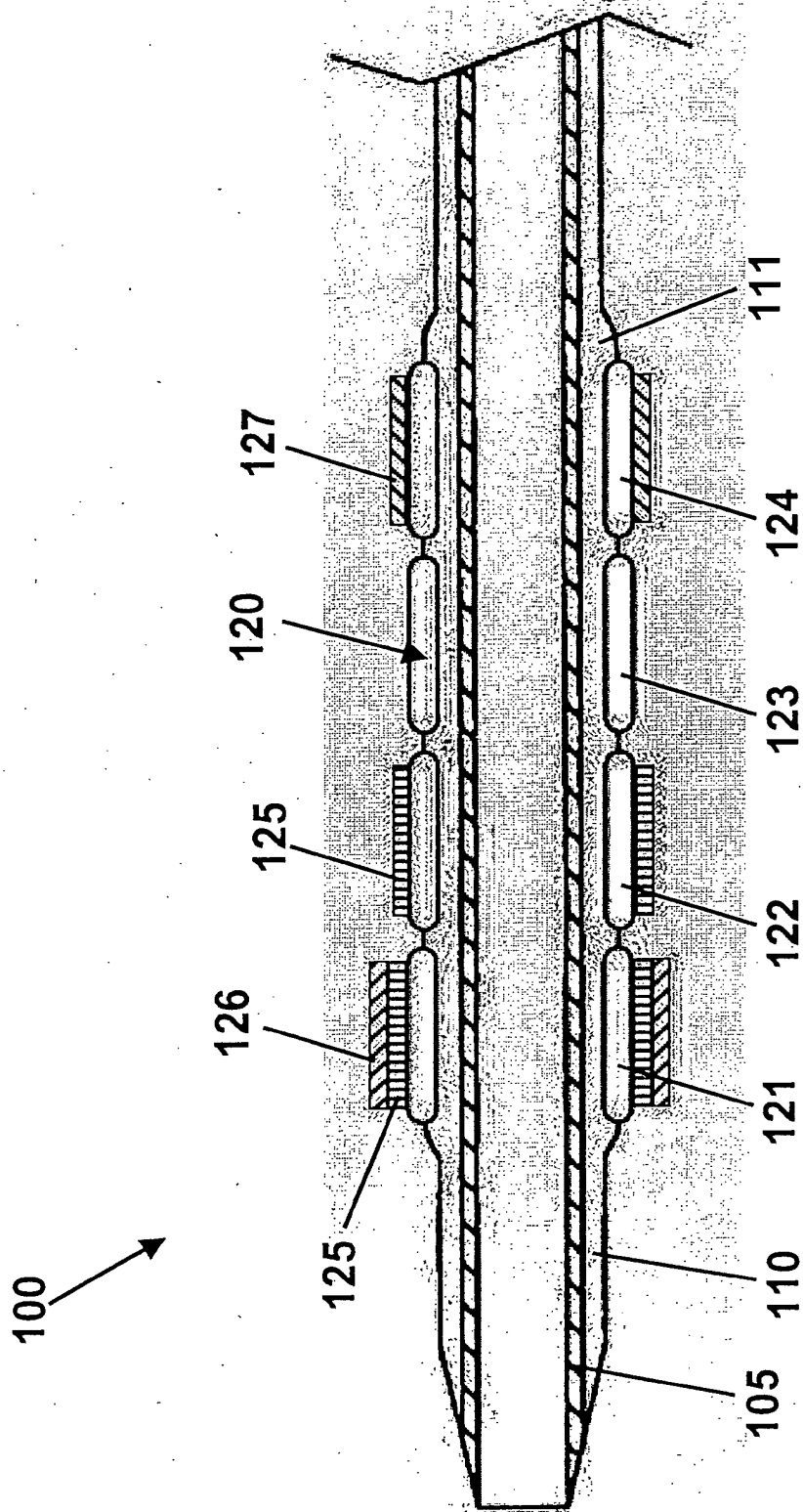


FIG. 1

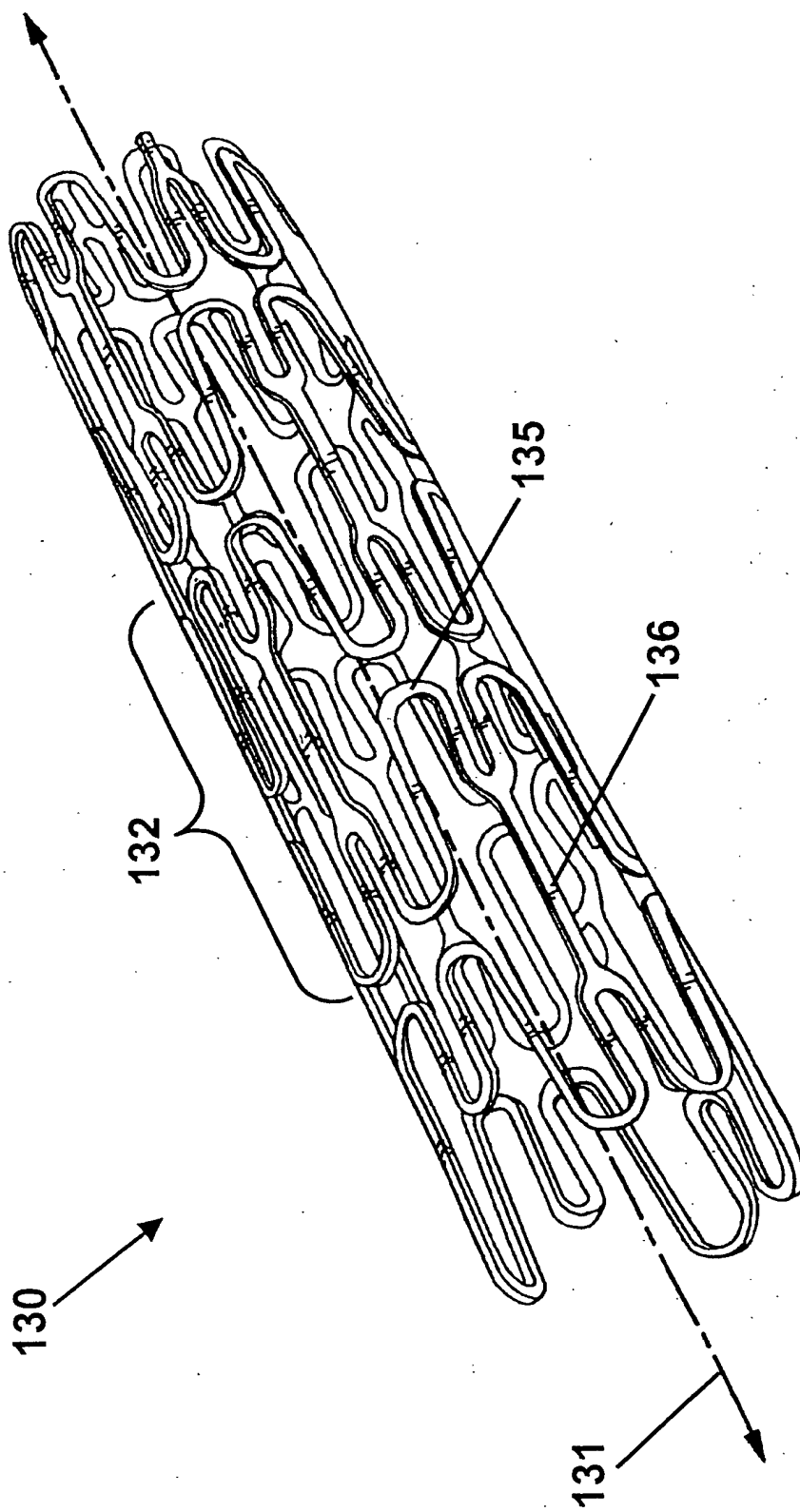
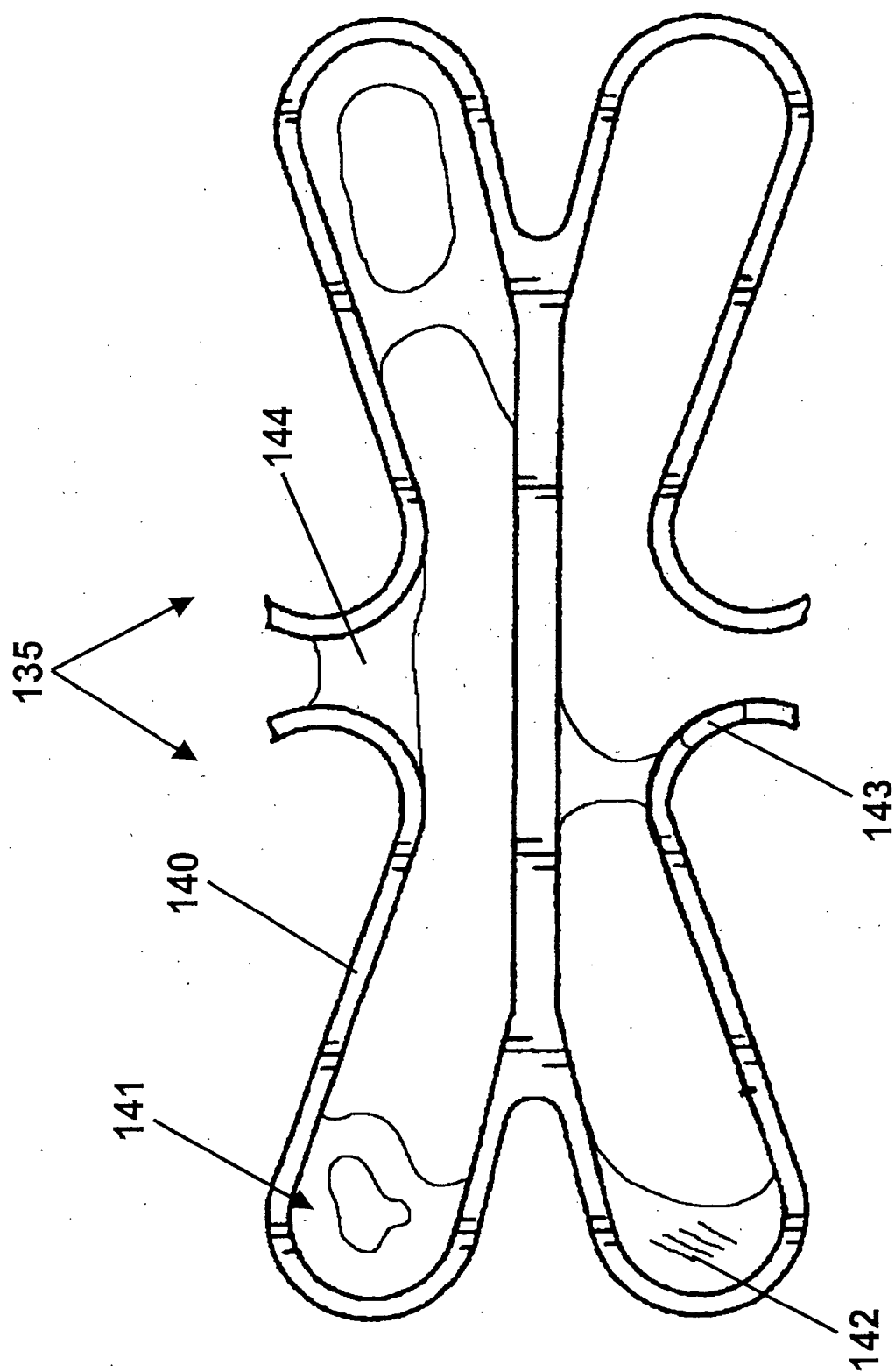


FIG. 2

**FIG. 3**

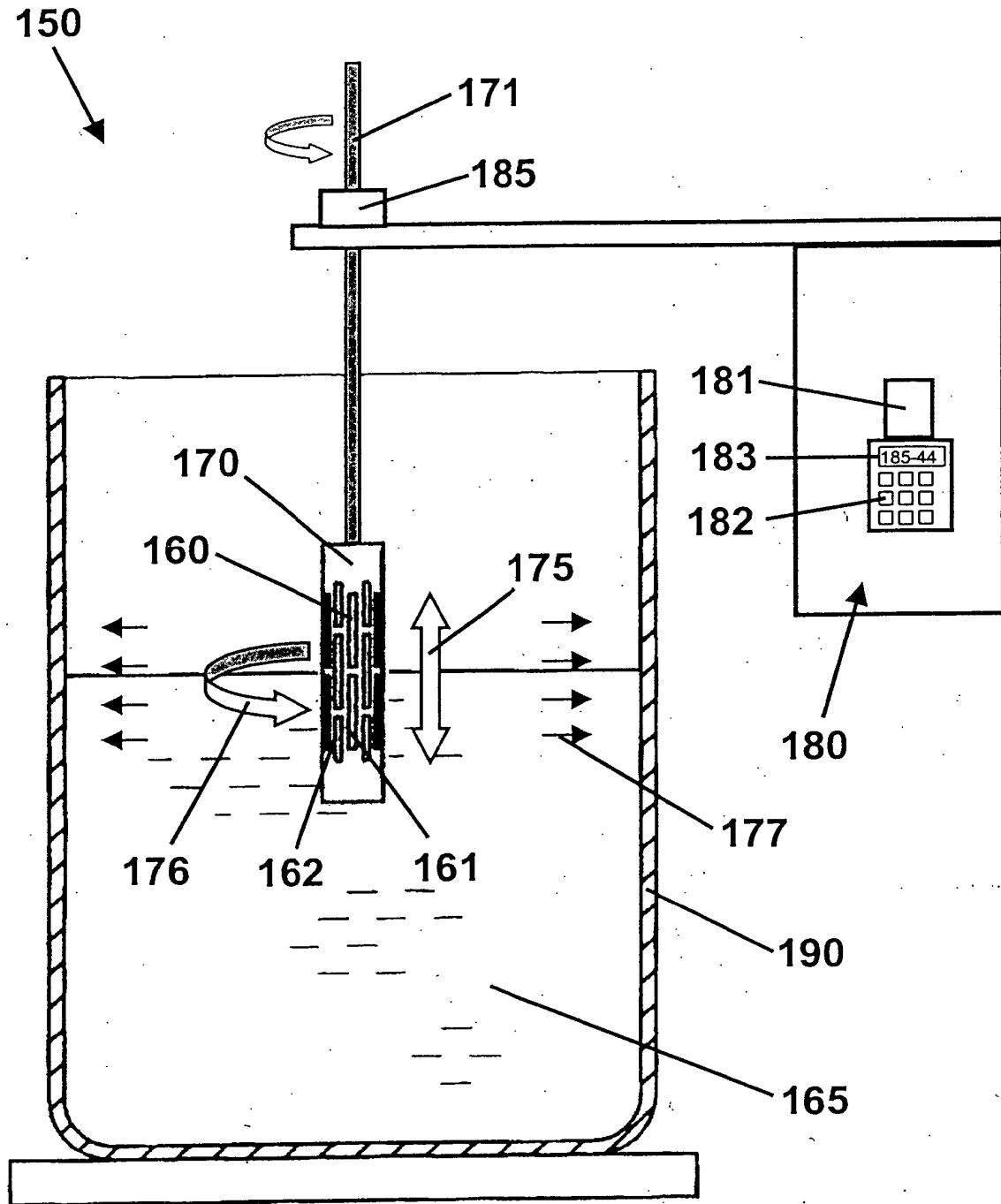


FIG. 4



European Patent  
Office

# EUROPEAN SEARCH REPORT

Application Number  
EP 03 00 0600

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
A	WO 00 17118 A (MANZANARES TAYLOR ELIZABETH R ; HARWOOD DUNCAN W J (GB); PAYNE DAVI) 30 March 2000 (2000-03-30) * claims; figures *	1-18	A61L31/10 B05D1/00 G03F7/16
A	WO 01 94103 A (RUEGG FRANK ARTHUR ; SOLA OPTICAL USA (US)) 13 December 2001 (2001-12-13) * claims *	1-18	
A	US 5 246 728 A (RODRIQUEZ JORGE M) 21 September 1993 (1993-09-21) * claims; examples *	1-18	
A	WO 00 29501 A (RADIOVASCULAR SYSTEMS L L C ; UNIV EMORY (US)) 25 May 2000 (2000-05-25) * claims; example 1 *	1-18	
A	EP 0 887 368 A (HUELS CHEMISCHE WERKE AG) 30 December 1998 (1998-12-30) * claims; example *	1-18	
			TECHNICAL FIELDS SEARCHED (Int.Cl.7)
			A61L B05D G03F
The present search report has been drawn up for all claims			
Place of search <b>THE HAGUE</b>		Date of completion of the search <b>26 May 2003</b>	Examiner <b>ESPINOSA, M</b>
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document			

EPO FORM 1503 03.92 (P04C01)

**ANNEX TO THE EUROPEAN SEARCH REPORT  
ON EUROPEAN PATENT APPLICATION NO.**

EP 03 00 0600

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on  
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

26-05-2003

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
WO 0017118	A	30-03-2000	AU	6102499 A	10-04-2000
			WO	0017118 A1	30-03-2000
WO 0194103	A	13-12-2001	AU	6457201 A	17-12-2001
			WO	0194103 A2	13-12-2001
US 5246728	A	21-09-1993	NONE		
WO 0029501	A	25-05-2000	AU	6495599 A	05-06-2000
			EP	1157076 A1	28-11-2001
			JP	2002530128 T	17-09-2002
			WO	0029501 A1	25-05-2000
			US	6475644 B1	05-11-2002
			US	2002077520 A1	20-06-2002
EP 0887368	A	30-12-1998	DE	19726735 A1	07-01-1999
			CA	2241381 A1	24-12-1998
			EP	0887368 A1	30-12-1998
			JP	11100429 A	13-04-1999
			NO	982942 A	28-12-1998

EPO FORM P0469

For more details about this annex : see Official Journal of the European Patent Office, No. 12/92